Amendments to the Claims:

This listing of claims will replace all prior versions, and listings of claims in the application.

Listing of Claims:

1. (Currently Amended) A compound of formula I

$$R_4$$
 X $(R_3)_n$ NH_2 (I) , R_1 N N

wherein

n is from 0 to 4,

R₁ is hydrogen, unsubstituted or substituted lower alkyl or halogen,

 R_2 is hydroxy; unsubstituted, mono- or disubstituted amino; an optionally substituted heterocyclic radical containing at least one nitrogen ring atom and being attached to the cyclohexane ring of the molecule of formula I via a nitrogen ring atom; a radical R_5 -(C=Y)-NH-, wherein R_5 is unsubstituted or substituted lower alkyl, unsubstituted, mono- or disubstituted amino, a heterocyclic radical, or etherified hydroxy, and Y is oxygen, sulfur or imino; or a radical R_6 -sulfonylamino, wherein R_6 is unsubstituted or substituted lower alkyl, unsubstituted, mono- or disubstituted amino or phenyl optionally substituted by lower alkyl, lower alkoxy or nitro, R_3 is lower alkyl, hydroxy-, amino- or halogen-substituted lower alkyl, hydroxy, cyano, lower alkoxy, lower alkanoyl, lower alkanoyloxy, amino, mono- or di-lower alkylamino, lower alkanoylamino, carboxy, lower alkoxycarbonyl or halogen, wherein the R_3 substituents can be selected independently of one another if n>1,

R₄ is a radical R₇-CR₈(R₉)-, wherein R₇ is cyclobutyl, cyclopentyl, cyclohexyl, phenyl, furyl, pyrrolyl, thienyl or pyridyl, said R₇ substitutents being optionally substituted by one or more radicals selected from lower alkyl and halogen, and R₈ and R₉ are independently of each other hydrogen, lower alkyl or halogen, and

X is selected from -O-, -NH- and -S-, or a salt thereof.

2. (Currently Amended) A compound of formula I according to claim 1, wherein n is from 0 to 4,

R₁ is hydrogen, unsubstituted or substituted lower alkyl or halogen,

R₂ is hydroxy; unsubstituted, mono- or disubstituted amino; an optionally substituted heterocyclic radical containing at least one nitrogen ring atom and being attached to the cyclohexane ring of the molecule of formula I via a nitrogen ring atom; a radical R₅-(C=Y)-NH-, wherein R₅ is unsubstituted or substituted lower alkyl, unsubstituted, mono- or disubstituted amino, a heterocyclic radical, or etherified hydroxy, and Y is oxygen, sulfur or imino; or a radical R₆-sulfonylamino, wherein R₆ is unsubstituted or substituted lower alkyl, unsubstituted, mono- or disubstituted amino or phenyl optionally substituted by lower alkyl, lower alkoxy or nitro, R₃ is lower alkyl or lower alkoxy, wherein the R₃ substituents can be selected independently of one another if n>1,

R₄ is a radical R₇-CR₈(R₉)-, wherein R₇ is cyclobutyl, cyclopentyl, cyclohexyl, phenyl, furyl, pyrrolyl, thienyl, pyridyl or phenyl substituted by one or more radicals selected from lower alkyl and halogen, and R₈ and R₉ are independently of each other hydrogen, lower alkyl or halogen, and

X is selected from -O-, -NH- and -S-, or a salt thereof.

R₁ is hydrogen, unsubstituted or substituted lower alkyl or halogen, R₂ is hydroxy; unsubstituted, mono- or disubstituted amino; an optionally substituted heterocyclic radical containing at least one nitrogen ring atom and being attached to the cyclohexane ring of the molecule of formula I via a nitrogen ring atom; a radical R₅-(C=Y)-NH-, wherein R₅ is unsubstituted or substituted lower alkyl unsubstituted mono- or disubstituted amino.

(Currently Amended) A compound of formula I according to claim 1, wherein is 0,

unsubstituted or substituted lower alkyl, unsubstituted, mono- or disubstituted amino, a heterocyclic radical, or etherified hydroxy, and Y is oxygen, sulfur or imino; or a radical R₆-sulfonylamino, wherein R₆ is unsubstituted or substituted lower alkyl, unsubstituted, mono- or disubstituted amino or phenyl optionally substituted by lower alkyl, lower alkoxy or nitro,

R4 is benzyl, and

3.

X is selected from -O-, -NH- and -S-, or a salt thereof.

4. (Currently Amended) A compound of formula I according to claim 1, wherein n is 0,

R₁ is hydrogen, unsubstituted or substituted lower alkyl or halogen,

R₂ is hydroxy; unsubstituted, mono- or disubstituted amino; an optionally substituted heterocyclic radical having from 4 to 8 ring members and from 1 to 3 heteroatoms whereby at least one heteroatom is nitrogen and the binding of the heterocyclic radical to the cyclohexane ring of the molecule of formula I occurs via a nitrogen ring atom; a radical R₅-(C=Y)-NH-, wherein R₅ is lower alkyl, unsubstituted, mono- or disubstituted amino, etherified hydroxy, a heterocyclic radical having from 4 to 8 ring members and from 1 to 3 heteroatoms whereby at least one heteroatom is nitrogen and the binding of the heterocyclic radical occurs via a nitrogen ring atom, lower alkyl substituted by said heterocyclic radical or by one or more radicals selected independently of one another from the group consisting of amino, N-lower alkylamino, N,N-dilower alkylamino, N-lower alkanoylamino, N,N-di-lower alkanoylamino, hydroxy, lower alkoxy, lower alkoxy-lower alkoxy, lower alkanoyl, lower alkanoyl, carbamoyl, amidino, guanidino, ureido, mercapto, lower alkylthio and halogen, and Y is oxygen, sulfur or imino; or a radical R₆-sulfonylamino, wherein R₆ is unsubstituted or substituted lower alkyl, unsubstituted, mono- or disubstituted amino or phenyl optionally substituted by lower alkyl, lower alkoxy or nitro,

R₄ is benzyl, and X is selected from -O-, -NH- and -S-, or a salt thereof.

5. (Currently Amended) A compound of formula I according to claim 1, wherein n is 0,

R₁ is hydrogen, lower alkyl or halogen,

R₂ is hydroxy; unsubstituted, mono- or disubstituted amino; an optionally substituted heterocyclic radical having from 4 to 8 ring members and from 1 to 3 heteroatoms whereby at least one heteroatom is nitrogen and the binding of the heterocyclic radical to the cyclohexane ring of the molecule of formula I occurs via a nitrogen ring atom; a radical R₅-(C=Y)-NH-, wherein R₅ is lower alkyl, unsubstituted or monosubstituted amino, etherified hydroxy, or lower alkyl substituted by a heterocyclic radical having from 4 to 8 ring members and from 1 to 3 heteroatoms whereby at least one heteroatom is nitrogen and the binding of the heterocyclic radical occurs via a nitrogen ring atom, and Y is oxygen or imino; or a radical R₆-sulfonylamino, wherein R₆ is lower alkyl or disubstituted amino,

R₄ is benzyl, and X is selected from -O-, -NH- and -S-, or a salt thereof.

6. (Original) A compound of formula I according to claim 1, wherein n is 0,

R₁ is hydrogen, lower alkyl or halogen,

 R_2 is hydroxy, amino, N,N-di-lower alkylamino, pyrimidinyl-amino, 1,4,5,6-tetrahydro-pyrimidinyl-amino, 4,5-dihydro-1H-imidazolyl-amino, azetidin-1-yl, pyrrolidin-1-yl, 1-piperidyl, lower alkyl-piperazin-1-yl, morpholin-4-yl, thiomorpholin-4-yl; a radical R_5 -(C=Y)-NH-, wherein R_5 is lower alkyl, lower alkoxy, amino, N-lower alkylamino, N-(phenyl-lower alkyl)-amino, N-(lower alkyl-phenyl-lower alkyl)-amino, N-(lower alkoxy-phenyl-lower alkyl)-amino, N-(morpholin-4-yl-lower alkyl)-amino, N-(N',N'-di-lower alkylamino-lower alkyl)-amino, lower alkoxy-lower alkoxy, 1-piperidyl-lower alkyl, morpholin-4-yl-lower alkyl or lower alkyl-piperazin-1-yl-lower alkyl, and Y is oxygen or imino; or a radical R_6 -sulfonylamino, wherein R_6 is lower alkyl or N,N-di-lower alkylamino,

R₄ is benzyl, and X is -O-, or a salt thereof.

7. (Original) A compound of formula I according to claim 1, selected from the group consisting of

cis-4-[4-amino-5-(3-benzyloxy-phenyl)-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexanol; trans-4-[4-amino-5-(3-benzyloxy-phenyl)-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexanol; cis-5-(3-benzyloxy-phenyl)-7-(4-piperidin-1-yl-cyclohexyl)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamine; trans-5-(3-benzyloxy-phenyl)-7-(4-piperidin-1-yl-cyclohexyl)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamine;

cis-5-(3-benzyloxy-phenyl)-7-(4-pyrrolidin-1-yl-cyclohexyl)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamine;

trans-5-(3-benzyloxy-phenyl)-7-(4-pyrrolidin-1-yl-cyclohexyl)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamine;

cis-5-(3-benzyloxy-phenyl)-7-[4-(4-methyl-piperazin-1-yl)-cyclohexyl]-7H-pyrrolo[2,3-d]pyrimidin-4-ylamine;

trans-5-(3-benzyloxy-phenyl)-7-[4-(4-methyl-piperazin-1-yl)-cyclohexyl]-7H-pyrrolo[2,3-d]pyrimidin-4-ylamine;

cis-5-(3-benzyloxy-phenyl)-7-(4-morpholin-4-yl-cyclohexyl)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamine;

trans-5-(3-benzyloxy-phenyl)-7-(4-morpholin-4-yl-cyclohexyl)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamine;

cis-7-(4-azetidin-1-yl-cyclohexyl)-5-(3-benzyloxy-phenyl)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamine; trans-7-(4-azetidin-1-yl-cyclohexyl)-5-(3-benzyloxy-phenyl)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamine;

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cis-5-(3-benzyloxy-phenyl)-7-(4-thiomorpholin-4-yl-cyclohexyl)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamine;
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trans-5-(3-benzyloxy-phenyl)-7-(4-thiomorpholin-4-yl-cyclohexyl)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamine;

trans-5-(3-benzyloxy-phenyl)-7-(4-diethylamino-cyclohexyl)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamine;

cis-7-(4-amino-cyclohexyl)-5-(3-benzyloxy-phenyl)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamine; trans-7-(4-amino-cyclohexyl)-5-(3-benzyloxy-phenyl)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamine; cis-{4-[4-amino-5-(3-benzyloxy-phenyl)-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexyl}-carbamic acid methyl ester;

cis-1-{4-[4-amino-5-(3-benzyloxy-phenyl)-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexyl}-3-methylurea;

cis-N-{4-[4-amino-5-(3-benzyloxy-phenyl)-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexyl}-2-piperidin-1-yl-acetamide;

cis-N-{4-[4-amino-5-(3-benzyloxy-phenyl)-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexyl}-2-morpholin-4-yl-acetamide;

cis-N-{4-[4-amino-5-(3-benzyloxy-phenyl)-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexyl}-2-(4-methyl-piperazin-1-yl)-acetamide;

cis-5-(3-benzyloxy-phenyl)-7-[4-(pyrimidin-2-ylamino)-cyclohexyl]-7H-pyrrolo[2,3-d]pyrimidin-4-ylamine;

cis-5-(3-benzyloxy-phenyl)-7-[4-(1,4,5,6-tetrahydro-pyrimidin-2-ylamino)-cyclohexyl]-7H-pyrrolo[2,3-d]pyrimidin-4-ylamine;

cis-5-(3-benzyloxy-phenyl)-7-[4-(4,5-dihydro-1H-imidazol-2-ylamino)-cyclohexyl]-7H-pyrrolo[2,3-d]pyrimidin-4-ylamine;

cis-N-{4-[4-amino-5-(3-benzyloxy-phenyl)-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexyl}-methanesulfonamide;

cis-N-{4-[4-amino-5-(3-benzyloxy-phenyl)-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexyl}-N,N-dimethylaminosulfonamide;

cis-5-(3-benzyloxy-phenyl)-7-(4-dimethylamino-cyclohexyl)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamine;

N-{4-[4-amino-5-(3-benzyloxy-phenyl)-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexyl}-acetamide; cis-1-{4-[4-amino-5-(3-benzyloxy-phenyl)-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexyl}-3-ethyl-urea; cis-1-{4-[4-amino-5-(3-benzyloxy-phenyl)-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexyl}-3-isopropyl-urea;

cis-1-{4-[4-amino-5-(3-benzyloxy-phenyl)-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexyl}-3-propylurea;

 $cis-1-\{4-[4-amino-5-(3-benzyloxy-phenyl)-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexyl\}-3-butyl-urea;\\$

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cis-1-{4-[4-amino-5-(3-benzyloxy-phenyl)-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexyl}-3-(3-methyl-
benzyl)-urea;
cis-1-{4-[4-amino-5-(3-benzyloxy-phenyl)-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexyl}-3-benzyl-urea;
cis-1-{4-[4-amino-5-(3-benzyloxy-phenyl)-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexyl}-3-(4-
methoxy-benzyl)-urea;
cis-1-{4-[4-amino-5-(3-benzyloxy-phenyl)-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexyl}-3-tert-butyl-
urea;
cis- N-{4-[4-amino-5-(3-benzyloxy-phenyl)-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexyl}-guanidine;
cis-1-[4-[4-amino-5-(3-benzyloxy-phenyl)-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexyl}-3-(2-
dimethylamino-ethyl)-urea;
cis-1-{4-[4-amino-5-(3-benzyloxy-phenyl)-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexyl}-3-(2-
morpholin-4-yl-ethyl)-urea;
cis-1-[4-[4-amino-5-(3-benzyloxy-phenyl)-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexyl}-3-(3-
morpholin-4-yl-propyl)-urea;
cis-[4-[4-amino-5-(3-benzyloxy-phenyl)-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexyl}-carbamic acid
2-methoxy-ethyl ester;
cis-4-[4-amino-5-(3-benzyloxy-phenyl)-6-bromo-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexanol;
trans-4-[4-amino-5-(3-benzyloxy-phenyl)-6-bromo-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexanol;
cis-4-[4-amino-5-(3-benzyloxy-phenyl)-6-methyl-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexanol;
trans-4-[4-amino-5-(3-benzyloxy-phenyl)-6-methyl-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexanol;
trans-5-(3-benzyloxy-phenyl)-6-methyl-7-[4-(4-methyl-piperazin-1-yl)-cyclohexyl]-7H-pyrrolo[2,3-
d]pyrimidin-4-ylamine;
trans-5-(3-benzyloxy-phenyl)-7-(4-dimethylamino-cyclohexyl)-6-methyl-7H-pyrrolo[2,3-
d]pyrimidin-4-ylamine;
trans-5-(3-benzyloxy-phenyl)-7-(4-diethylamino-cyclohexyl)-6-methyl-7H-pyrrolo[2,3-
d]pyrimidin-4-ylamine;
trans-5-(3-benzyloxy-phenyl)-6-methyl-7-(4-pyrrolidin-1-yl-cyclohexyl)-7H-pyrrolo[2,3-
d]pyrimidin-4-ylamine;
trans-5-(3-benzyloxy-phenyl)-6-methyl-7-(4-morpholin-4-yl-cyclohexyl)-7H-pyrrolo[2,3-
d]pyrimidin-4-ylamine;
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8. (Cancelled) A compound of formula I, or a pharmaceutically acceptable salt thereof,

4-ylamine;

and pharmaœutically acceptable salts thereof.

trans-7-(4-azetidin-1-yl-cyclohexyl)-5-(3-benzyloxy-phenyl)-6-methyl-7H-pyrrolo[2,3-d]pyrimidin-

according to claim 1 for use in a method for the treatment of the human or animal body.

9. (Previously Presented) A pharmaceutical composition comprising a compound of formula I or a pharmaceutically acceptable salt thereof according to claim 1, together with at least one pharmaceutically acceptable carrier.

10.-12. (Cancelled)